

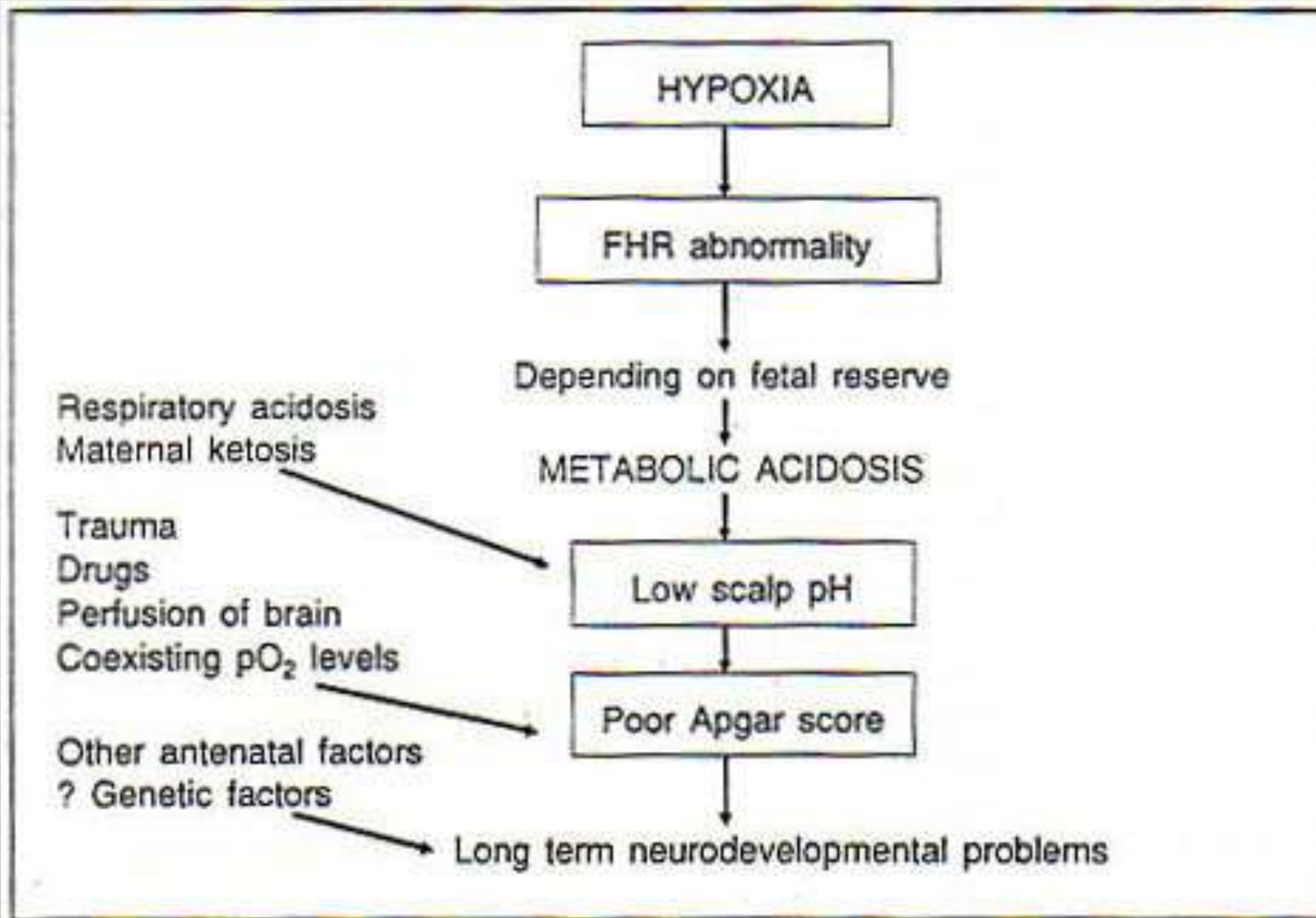


Intrapartum fetal surveillance

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Intrapartum fetal surveillance

- The aim is to detect potential fetal decompensation and to allow timely and effective intervention to prevent perinatal/neonatal morbidity or mortality.
- Changes in fetal heart rate precede brain injury, so timely response to abnormal fetal heart patterns might be effective in preventing brain injury.
- During uterine contractions there is a decrease in uteroplacental blood flow and a subsequent increase in fetal pCO₂ and a decrease in pO₂ and PH.
- Avoiding adverse fetal outcome related to hypoxia/acidosis is the main objective of intrapartum fetal monitoring.



The relationship between hypoxia, intrapartum fetal monitoring and neonatal outcome.

Antenatal factors that increase the risk of fetal compromise

- Oligohydramnios or polyhydramnios
- Multiple pregnancy
- Antepartum hemorrhage
- Previous caesarean section
- Hypertension or pre-eclampsia and diabetes
- Prolonged pregnancy
- Intrauterine growth restriction

Antenatal factors that increase the risk of fetal compromise

- Induction of labour with prostaglandin/oxytocin
- Regional anesthesia
- Maternal pyrexia: $\geq 38^{\circ}\text{C}$
- Meconium or blood stained liquor
- Pre-term labour
- Uterine hyperstimulation

Antenatal factors that increase the risk of fetal compromise

- Contractions last longer than 2 minutes, or 5 or more contractions in 10 minutes.
- Presence meconium.
- Maternal pyrexia (a temperature of 38°C or above on a single reading or 37.5°C or above on 2 consecutive occasions 1 hour apart).
- Suspected chorioamnionitis or sepsis.
- Fresh vaginal bleeding that develops in labor, or blood-stained liquor .
- Maternal pulse over 120 beats a minute on 2 occasions 30 minutes apart.
- Confirmed delay in the first or second stage of labor

Intrapartum fetal surveillance

- **Fetal Heart Rate Monitoring (FHR)**
 - Intermittent auscultation
 - Continuous electronic fetal heart monitoring (EFM)
- **Fetal Scalp blood pH estimation**
- **Intrapartum fetal stimulation tests**
 - Fetal scalp stimulation tests
 - Fetal acoustic stimulation test (FAST)
- **Newer approaches**
 - Fetal ECG
 - Scalp blood lactate estimation
 - Continuous biochemical monitoring
(Pulse oximetry)

Intermittent Auscultation

- It is the recommended fetal surveillance method during labor for healthy women without risk factors for adverse perinatal outcome.
- A baseline heart rate is assessed by listening and counting FHR between uterine contractions.
- FHR is counted for **60 seconds**.

Intermittent Auscultation

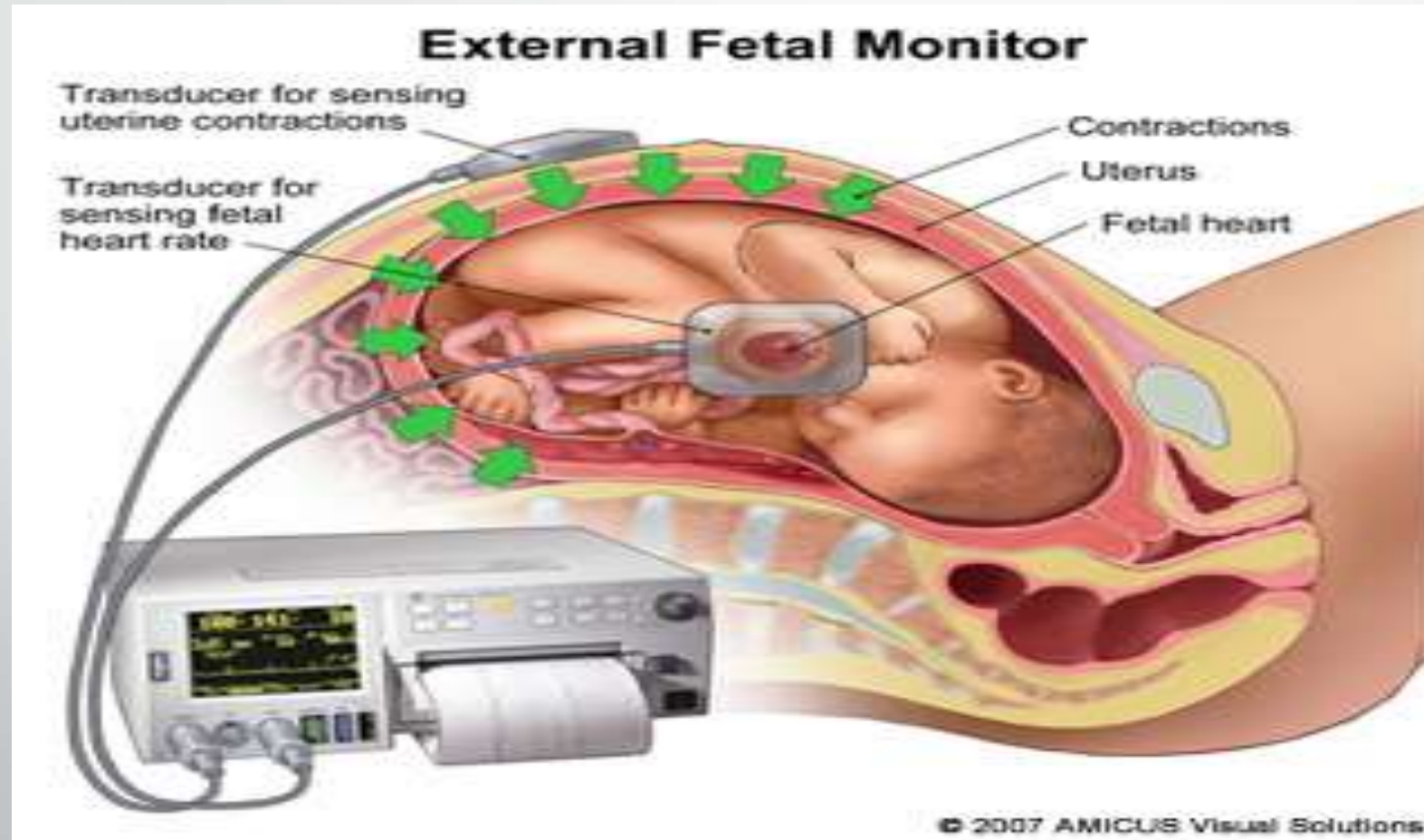
- The FHR should be assessed at least every **15 minutes** in the first stage of labor and every **5 minutes** in the second stage .
- Palpate the woman's pulse simultaneously to differentiate between the maternal and fetal heart rates.
- It ensures frequent contact between healthcare professionals and the laboring woman.
- If abnormal, EFM is recommended .

Pinard stethoscope



Handheld Doppler device

Electronic fetal heart monitoring



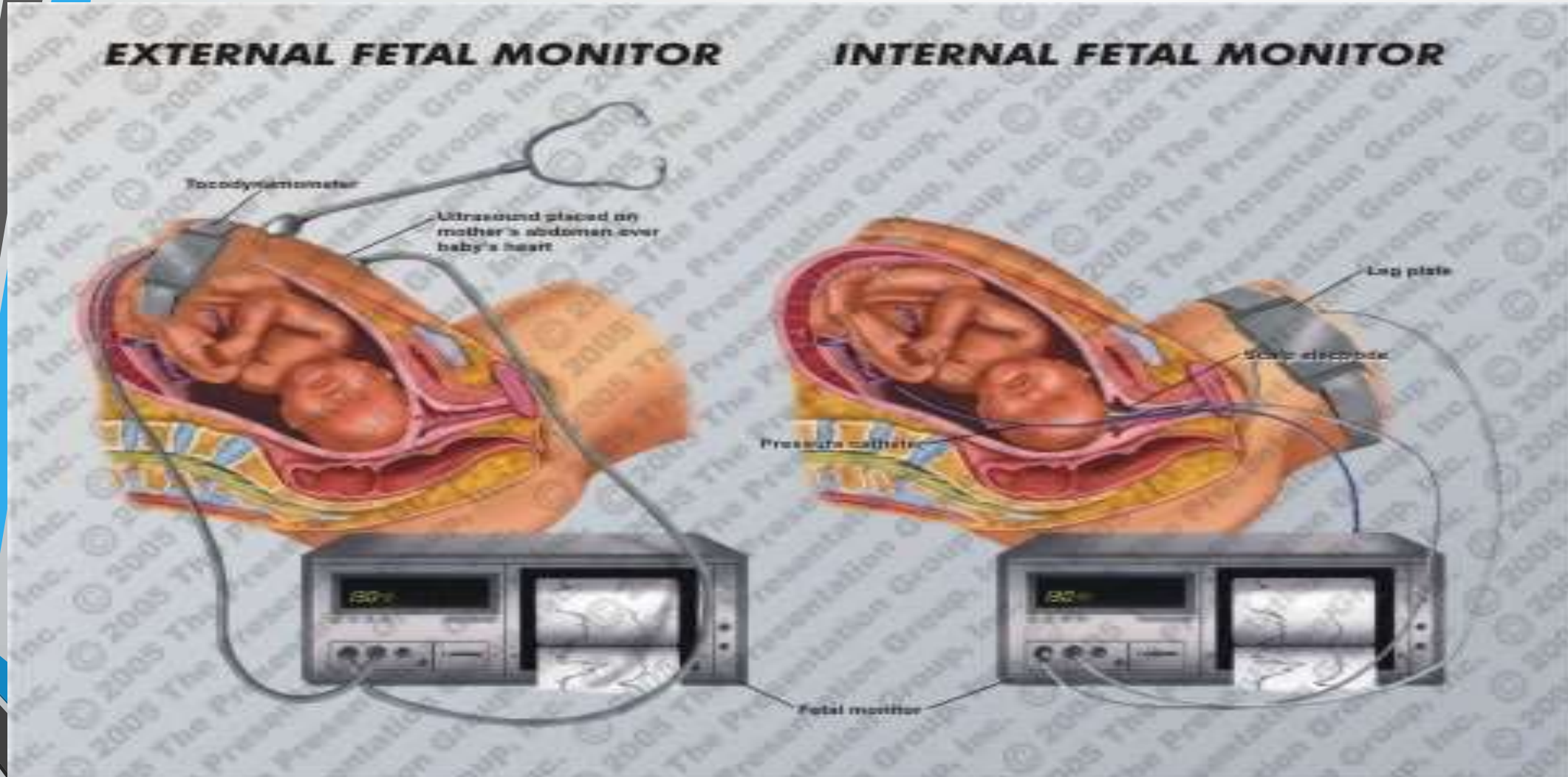
Electronic fetal heart monitoring

- It should be considered in all situations where there is a high risk of fetal hypoxia/acidosis.
- The electronic FHR monitor is a device with two components. One establishes the FHR, and the other measures uterine contractions.
- Continuous cardiotocography (CTG)is also recommended when abnormalities are detected during intermittent fetal auscultation.
- CTG has been shown to decrease the occurrence of neonatal seizures.

Electronic fetal heart monitoring

- Limit woman's mobility
- Decrease direct contact between woman & staff.
- Continuous EFM is associated with an increase in the rates of Caesarean sections and instrumental vaginal births.

EFM may be performed with an external or internal monitor



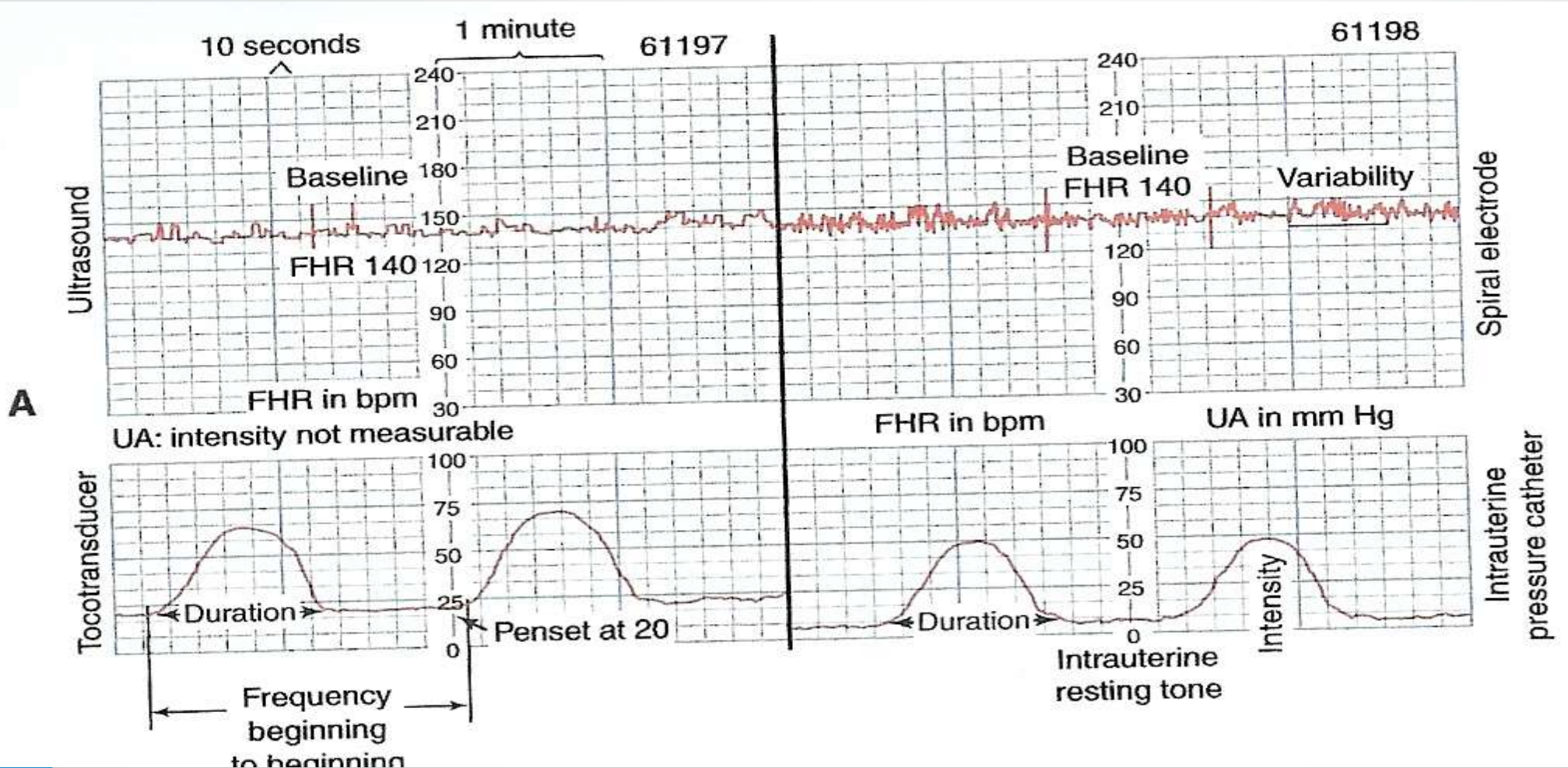
External fetal monitor

- An ultrasound transducer transmits the FHR in beats per minute (bpm).
- Non-invasive.
- Does not require cervical dilatation or rupture of membranes.
- Needs readjustment with maternal or fetal movements.
- Difficult to obtain a clear tracing in obese women or those with polyhydramnios.

Internal fetal monitor

- Performed with a spiral electrode inserted through vagina and cervix and attached to the fetal scalp.
- Indicated when the external tracing is inadequate for accurate interpretation.
- Contraindications include placenta previa, face presentation, unknown presentation, HIV seropositivity, or active genital herpes.
- Internal uterine activity monitoring is done via an IUPC.

External vs internal monitoring



Electronic fetal heart monitoring

- Baseline FHR
- FHR variability (beat to beat variation)
- Acceleration
- Deceleration

Interpretation of electronic fetal monitoring

1. Assess the quality of the signal acquisition.
2. Determine the paper speed and graph range.
3. Determine whether the mode of recording is external or internal.
4. Assess the uterine activity pattern, including frequency, duration, and intensity of contraction, and uterine resting tone.
5. Assess the FHR.

Assessment of uterine contractions

- The lowest intrauterine pressure between contractions is called resting tone.
- Normal resting tone is 5-10 mmHg, but during labor it may rise to 10-15 mmHg.
- Pressure during contractions rises to ~25-100 mmHg (varies with stage).
- A resting pressure above 20 mmHg causes decreased uterine perfusion.

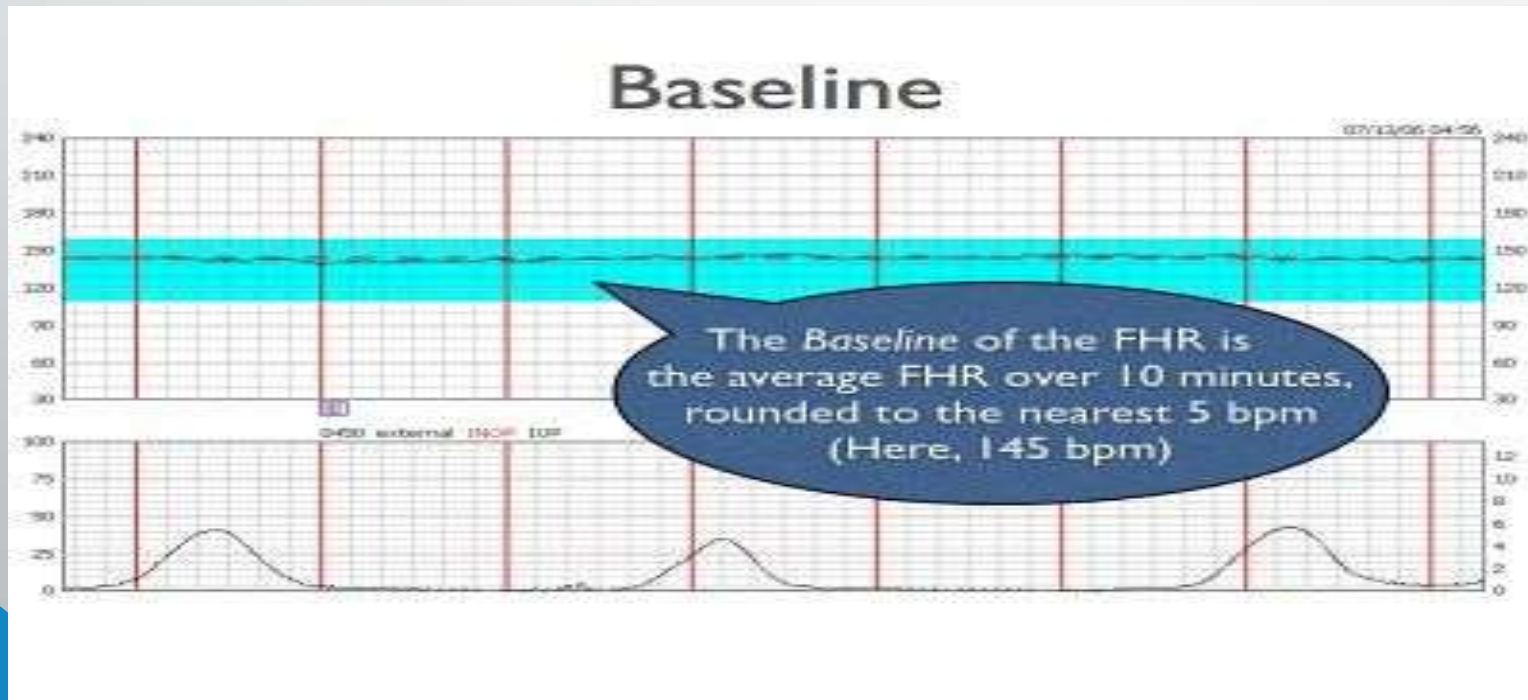
Assessment of uterine contractions

- White: fewer than 5 contractions in 10 minutes.
- Amber: 5 or more contractions in 10 minutes, leading to reduced resting time between contractions, or hypertonus.

Assessment of fetal heart rate

- **The baseline FHR :**

- Baseline FHR is the mean FHR rounded to 5bpm, excluding accelerations and decelerations over a period of 10 minutes .
- Normal baseline rate is between 110 and 160 bpm .

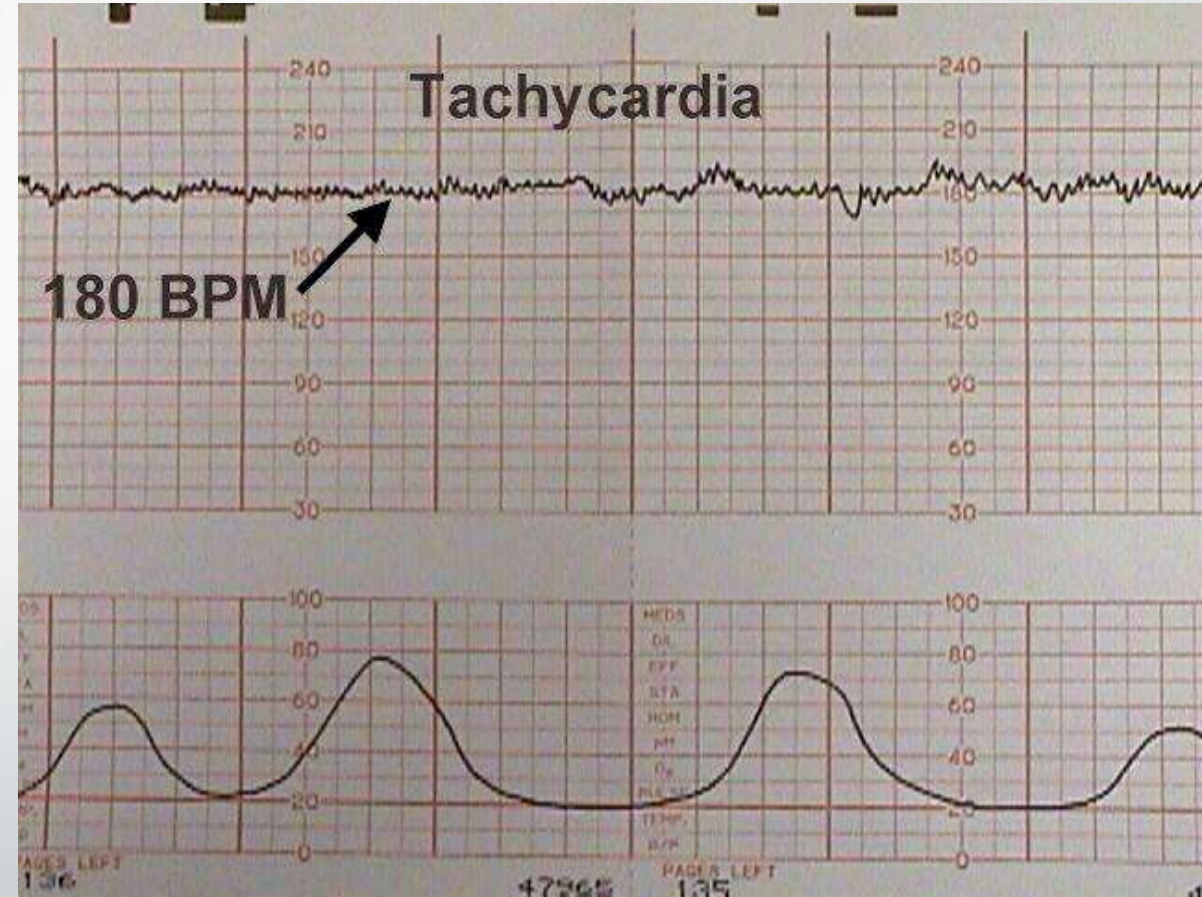


Tachycardia

- A baseline value above 160 bpm lasting more than 10 minutes.
- Maternal pyrexia (infection, epidural analgesia) is the most frequent cause of fetal tachycardia.
- Other causes
 - Fetal hypoxia.
 - Medications (beta-agonist drugs) .
 - Fetal arrhythmias (SVT).
 - Fetal anemia.

Tachycardia

- Left lateral position
- IV hydration
- Oxygen
- Stop oxytocin

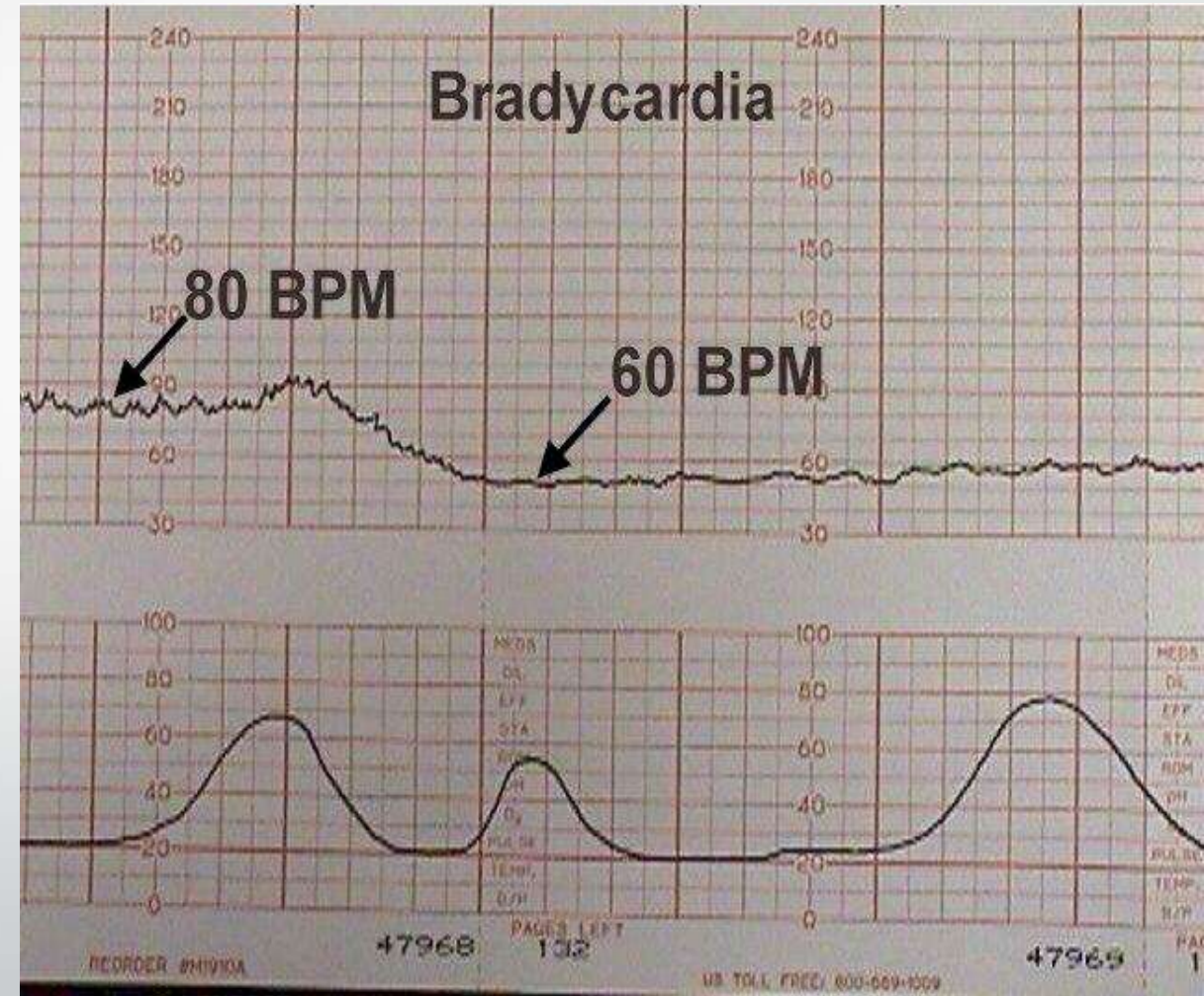


Bradycardia

- A baseline value below 110 bpm lasting more than 10 minutes.
- Values between 100 and 110 bpm may occur in normal fetuses, especially in postterm pregnancies.
- Sudden drop in oxygenation, such as placental abruption.^{[L][SEP]}
- Decrease or cessation in umbilical blood flow, such as occurs with a prolapsed cord or uterine rupture.
- Maternal hypothermia, maternal hypotension, administration of beta-blockers, and fetal arrhythmias such as atrioventricular block are other possible causes.

Bradycardia

- Left lateral position.
- Increase IV hydration.
- Oxygen
- Vaginal exam



The baseline FHR

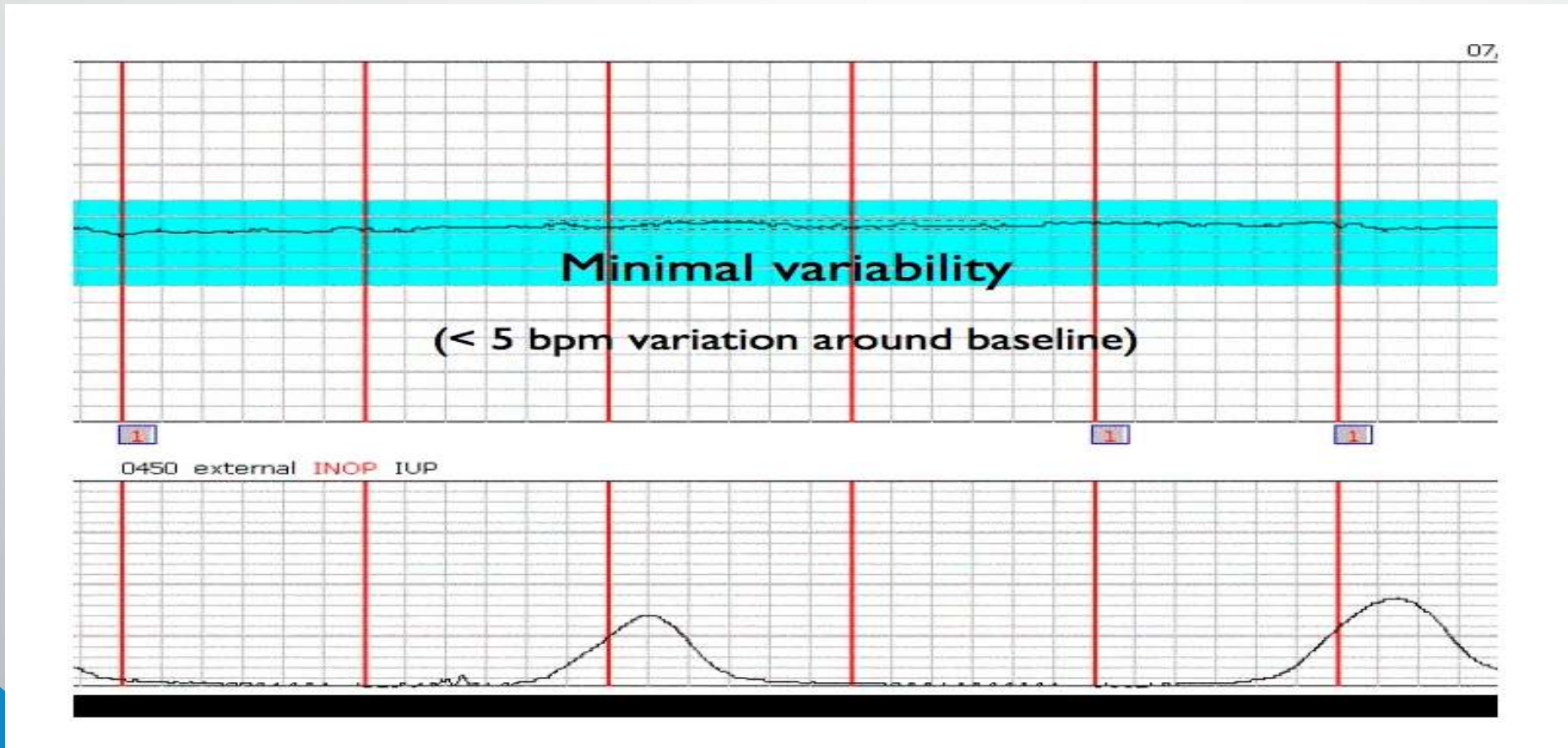
- **White:** stable baseline of 110 to 160 beats a minute.
- **Amber:** increase in baseline fetal heart rate of 20 beats a minute or more from the start of labor, or: 100 to 109 beats a minute.
- **Red:** below 100 beats a minute, or above 160 beats a minute.

Assessment of the FHR

- **baseline variability:**
 - It refers to the fluctuations in the baseline FHR. (minor oscillations in the fetal heart rate)
 - Measure it by estimating the difference in beats per minute between the highest heart rate and the lowest heart rate in a 1-minute segment of the trace between contractions, excluding decelerations and accelerations.
 - Normal variability of 5–25 bpm.
 - Hypoxia and acidosis, fetal sleep, medications, (e.g., narcotics, sedatives, b-blockers, betamethasone), prematurity, fetal tachycardia, and congenital anomalies decrease FHR variability.

Assessment of the FHR

- Less than 5 bpm minimal
- 5 to 25 bpm moderate
- > 25 bpm marked



Assessment of the FHR

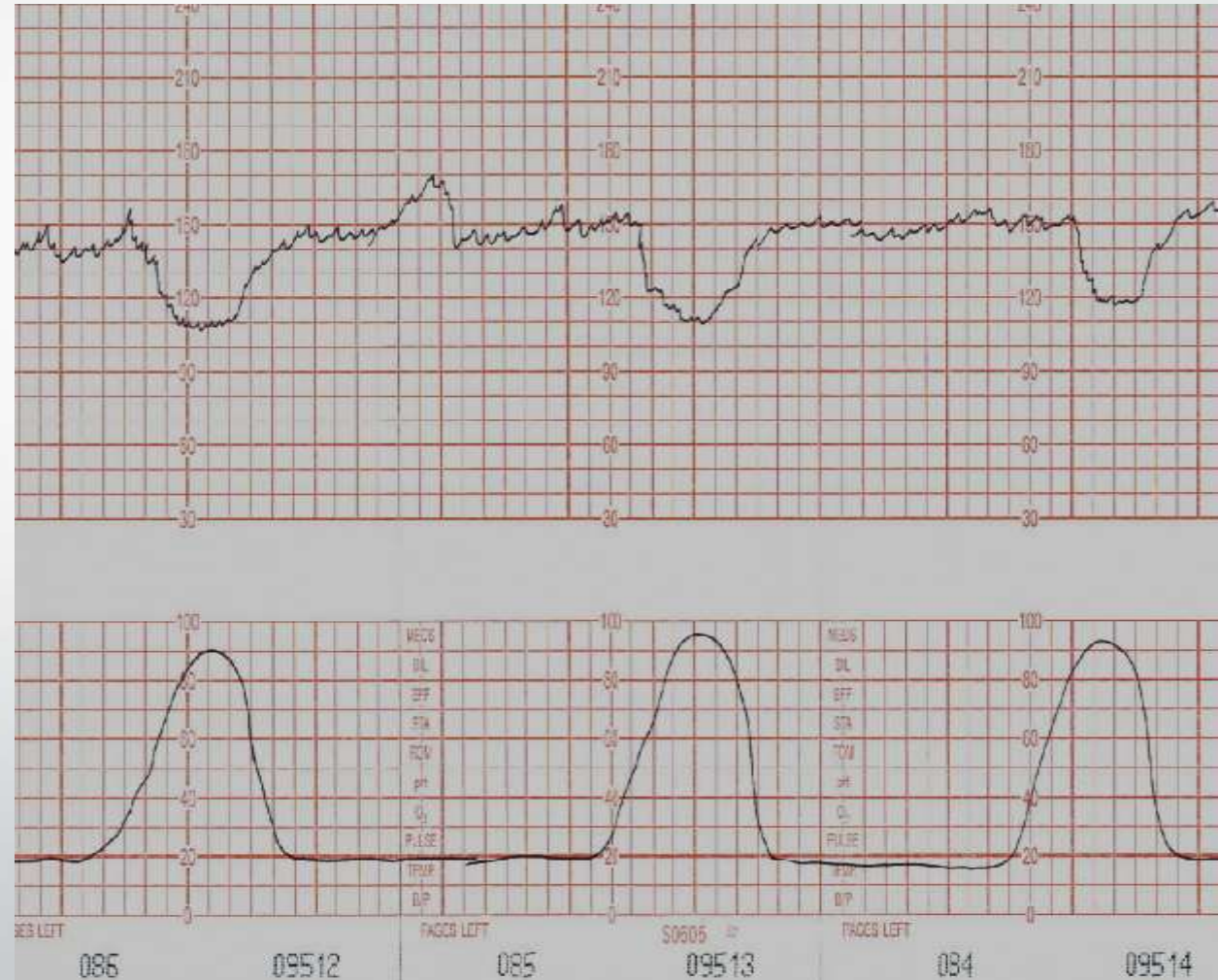
- **baseline variability:**
- **White:** 5 to 25 beats a minute
- **Amber:**
 - Less than 5 beats a minute for between 30 and 50 minutes.
 - More than 25 beats a minute for up to 10 minutes.
- **Red:**
 - less than 5 beats a minute for more than 50 minutes.
 - More than 25 beats a minute for more than 10 minutes.
 - Sinusoidal.

Assessment of the FHR

- **Decelerations:**
 - Transient episodes when the FHR slows to below the baseline level by more than **15 beats a minute**, with each episode lasting **15 seconds or more**.
 - **Their timing** (**early, variable or late**) in relation to the peaks and duration of the contractions
 - **The duration** of the individual decelerations
 - The FHR **returns to the baseline**.
 - **How long** they have been present
 - Occur with **over 50% of contractions** (repetitive)
 - Presence or absence of **shouldering**
 - **Variability** within the deceleration

Early decelerations

- gradual decrease in the FHR and return to baseline associated with uterine contraction.
- The onset, nadir, and recovery of the decelerations coincide with the beginning, peak, and ending of the contraction.
- Caused by fetal head compression and do not indicate fetal hypoxia/acidosis.



Variable decelerations

▼ = 15 bpm/15 secs

Variable Deceleration

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Abrupt onset to nadir < 30 secs, with drop of 15 bpm below baseline for ≥ 15 secs but < 2 min

Onset Nadir Recovery

Baseline

Onset to nadir < 30 secs

1250 FECG INOP INOP TOCO

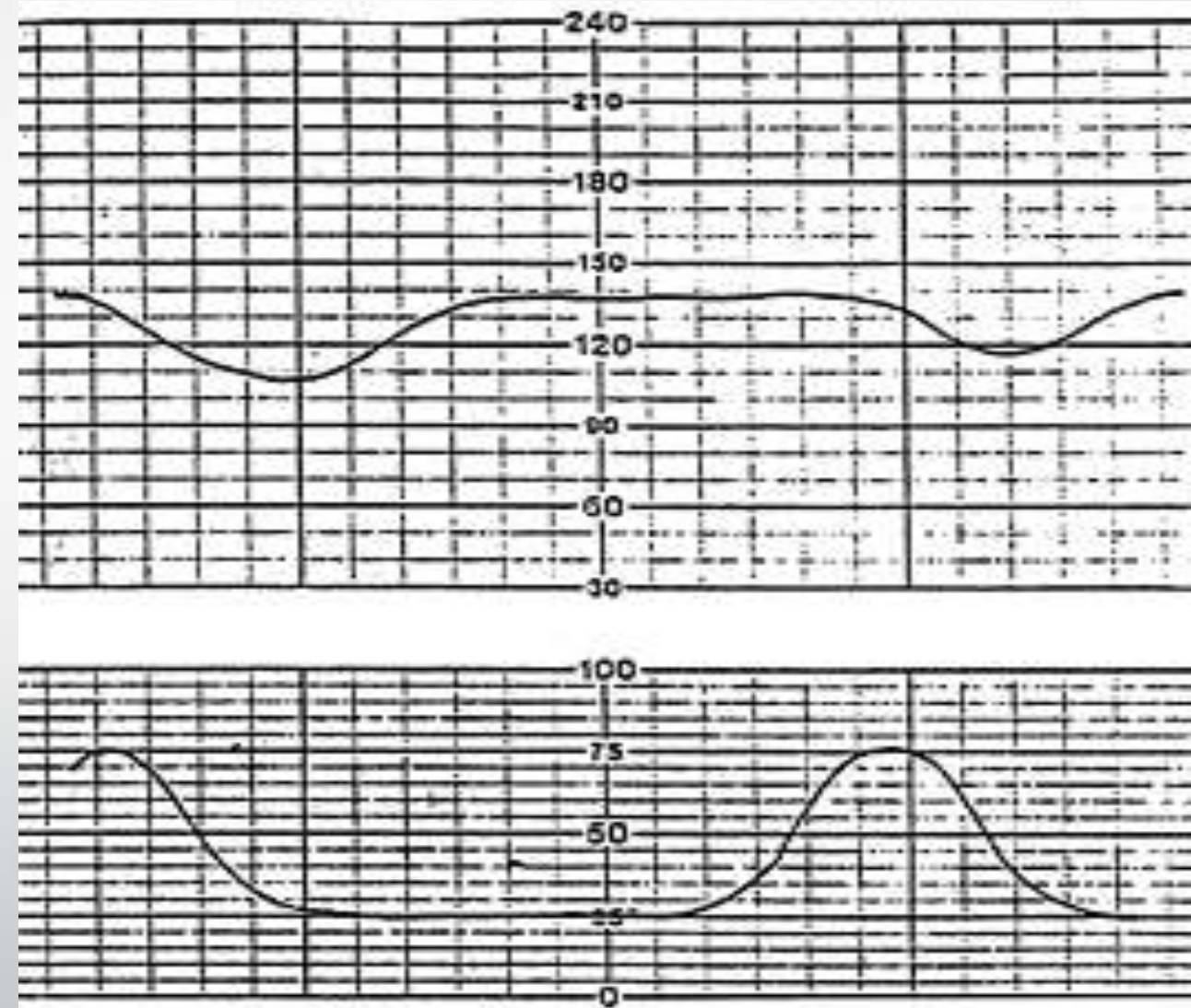
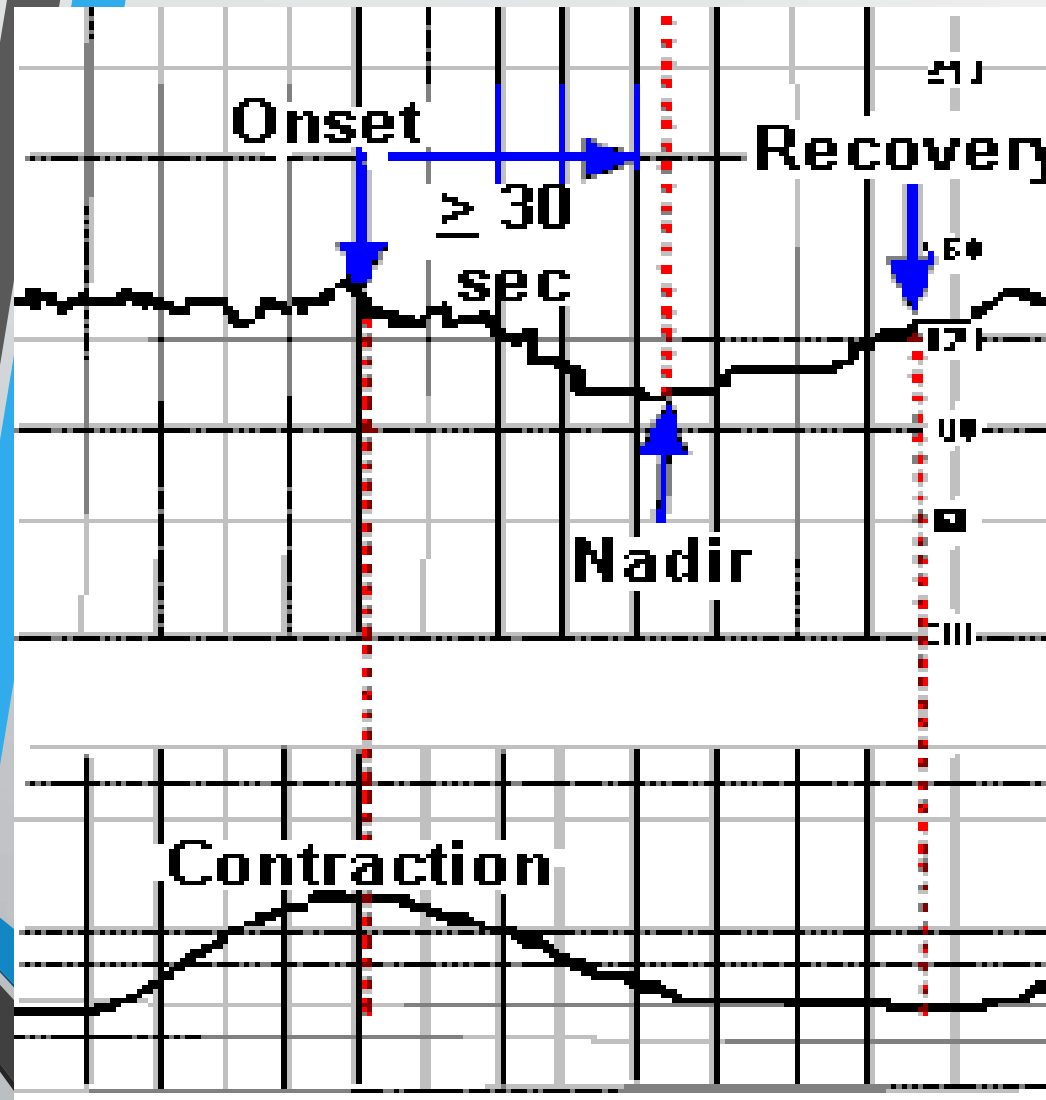
The relationship to the contraction is variable

- Most common type
- Caused by chemoreceptor stimulation secondary to cord compression

Variable decelerations

- Concerning characteristics of variable decelerations:
 - Lasting more than 60 seconds.
 - Reduced variability within the deceleration.
 - Failure or slow return to baseline fetal heart rate.
 - Loss of previously present shouldering.

Late decelerations



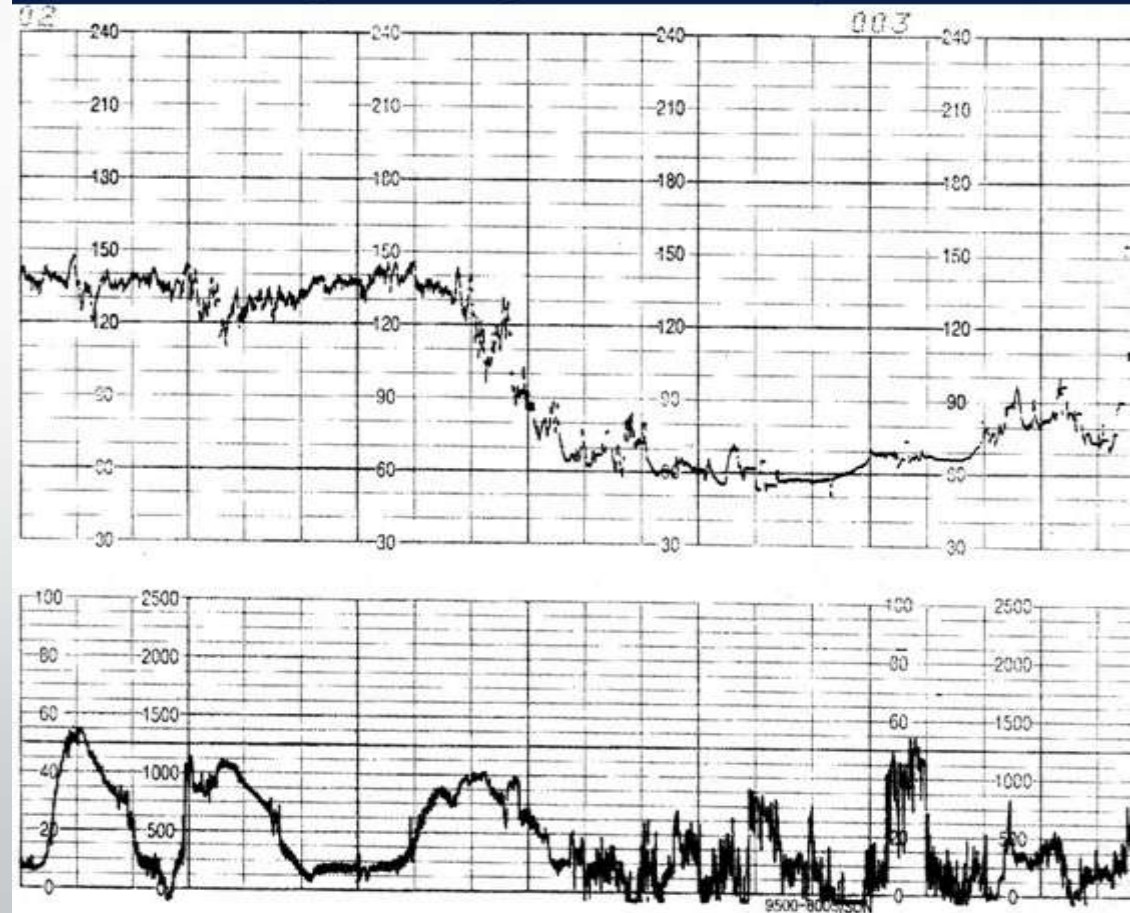
Late decelerations

- Late decelerations are found in association with uteroplacental insufficiency.
- Management :
 - Maternal left lateral position
 - Correct maternal hypotension with IV fluids
 - Stop oxytocin infusion
 - Administer O2 by mask
 - Vaginal examination
 - If persistent perform fetal scalp PH

Prolonged decelerations

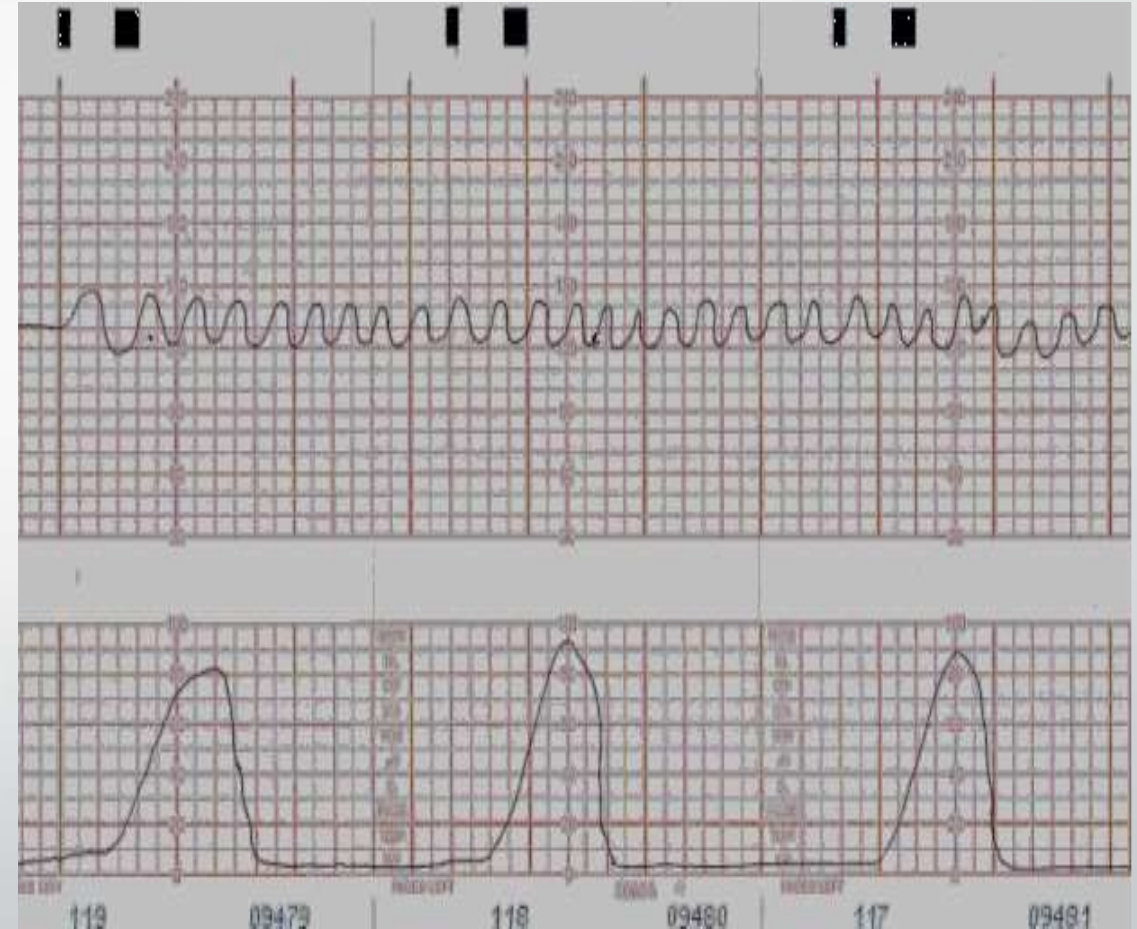
- Usually lasting more than 3 minutes.
- They indicate hypoxia
- If associated with reduced variability, they indicate acute fetal hypoxia/acidosis and require emergent intervention.

Fig 6 Prolonged Deceleration



Sinusoidal pattern

- A regular, smooth, undulating signal, resembling a sine wave.
- It occurs in association with severe fetal anemia (anti-D alloimmunization, fetal-maternal hemorrhage, and ruptured vasa previa)



Decelerations

- **White:**
 - No decelerations.
 - Early decelerations.
 - Variable decelerations with no concerning characteristics.

Decelerations

- Amber:
 - Repetitive variable decelerations with any concerning characteristics for < 30 minutes.
 - Variable decelerations with any concerning characteristics for > 30 minutes.
 - Repetitive late decelerations for <30 minutes .

Decelerations

- Red:
 - Repetitive variable decelerations with any concerning characteristics for > 30 minutes.
 - Repetitive late decelerations for > 30 minutes.
 - Single prolonged deceleration lasting 3 minutes or more.

Interpretation of FHR

Interpretation	Baseline (bpm)	Variability (bpm)	Decelerations	Accelerations
White (Reassuring)	110-160	5-25	<ul style="list-style-type: none"> - No decelerations - Early decelerations - 	present
Amber (Non-reassuring)	161-180 100- 109	<ul style="list-style-type: none"> - < 5 for 30–50 minutes - > 25 up to 10 minutes 	<ul style="list-style-type: none"> - Repetitive Variable decelerations < 30 minutes. - Variable decelerations with concerning features > 30 minutes. - Repetitive Late decelerations, < 30 minutes. 	
Red (Abnormal)	Above 180 Below 100	<ul style="list-style-type: none"> - < 5 for over 50 minutes - > 25 for > 10 minutes - Sinusoidal 	<ul style="list-style-type: none"> - Repetitive Variable decelerations > 30 minutes. - Repetitive Late decelerations > 30 minutes. - Single prolonged deceleration more than 3 min. 	

Interpretation of FHR

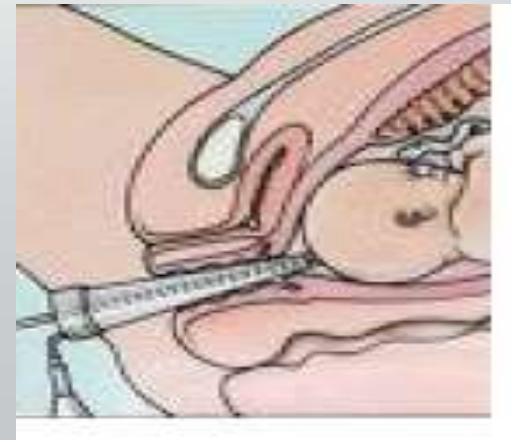
Category	Definition	Management
CTG is normal/ reassuring	no amber or red features	Continue CTG and normal care.
CTG is suspecious	1 amber feature, and 2 white features	<ul style="list-style-type: none">- check temperature- left-lateral position- oral or intravenous fluids- stopping oxytocin- Inform coordinating midwife and obstetrician- Scalp stimulation
CTG is abnormal and indicates further testing	1 red feature OR 2 amber features	<ul style="list-style-type: none">- Same as non reassuring- Consider urgent delivery

Secondary tests of Fetal wellbeing

- Fetal Scalp Sampling
- Scalp stimulation
- Acoustic stimulation
- Fetal pulse oximetry
- Fetal Electrocardiogram Analysis

Fetal blood sampling for pH and lactate

- FBS may be used in cases of abnormal CTG.
- A vaginal examination needs to be performed prior to the procedure to assess the nature and position of the presenting part.
- Contraindications : maternal infection, women seropositive to hepatitis B, C, or to HIV, suspected fetal blood disorders, uncertainty about the presenting part, preterm fetus.
- FBS (lack of evidence to support doing it)



Fetal blood sampling for pH and lactate

Interpretation	pH	Lactate (mmol/L)
Normal	≥ 7.25	< 4.2
Repeat in 30 mins	7.21 – 7.24	4.2 – 4.8
Birth expedited	≤ 7.20	> 4.8
Urgent delivery	< 7.15	> 5.0

Fetal stimulation tests

Test	Recommendation
Digital stimulation	Digital stimulation of the fetal scalp during vaginal exam may be considered as an adjunct to FHM
Vibroacoustic stimulation	Of value in non-reactive NST, but no prove in assessment during labour
Maternal glucose ingestion	No evidence to improve fetal wellbeing
Manual fetal manipulation	This procedure is not recommended

Fetal pulse oximetry

- Monitor intrapartum fetal O₂ saturation
- Fetal pulse oximetry is a relatively new technique in the assessment of a fetus prior to delivery
- It measures both the pulse rate and oxyhemoglobin saturation
- Sensor is placed transvaginally through the cervix to rest against the fetal cheek or temple, requiring cervical dilatation (~ 2 cm or more) and ruptured amniotic membranes with a cephalic presentation.
- Insufficient evidence to substantiate a recommendation for the use of Fetal Pulse oximeter as an adjunct or independent of electronic fetal surveillance.

Fetal Electrocardiogram Analysis

- Used in combination with standard EFM.
- Specialized monitor with software collects both the familiar fetal heart rate and uterine activity signals, and the fetal ECG
- Interpretation is based on the observation that the fetal QRS and T wave change in relation to the metabolic state of the fetal heart
- The impact of this type of monitoring compared with standard EFM was studied and showed no difference in the number of Caesarean sections, perinatal deaths, or NICU admissions.